

X. SUMMARY OF SAFETY AND EFFECTIVENESS**JAN 0 8 2002**

SUBMITTER: United States Surgical,
a division of Tyco Healthcare Group, LP
150 Glover Avenue
Norwalk, CT 06856

CONTACT PERSON: Robert Zott

DATE PREPARED: November 1, 2001

CLASSIFICATION NAME: Absorbable poly(glycolide/L-lactide)
surgical suture

COMMON NAME: Surgical Suture

PROPRIETARY NAME: To be determined

PREDICATE DEVICES: K885018: Gut Suture (Surgigut™)
K000037: Biosyn™ Suture
K944110: Vicryl™ Rapide Suture

INDICATIONS: The device is indicated for use in general soft
tissue approximation and/or ligation, but not for
use in cardiovascular or neurological surgery,
microsurgery, or ophthalmic surgery.

MATERIALS: Glycolide, Caprolactone, Tri-Methylene
Carbonate, and Lactide polymers.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 0 8 2002

Robert Zott
Senior Associate, Regulatory Affairs
United States Surgical
150 Glover Avenue
Norwalk, Connecticut 06856

Re: K013671
Trade Name: L-25 Absorbable Surgical Suture
Regulation Number: 878.4493
Regulation Name: Absorbable polyglycolide surgical suture
Regulatory Class: II
Product Code: GAM
Dated: November 5, 2001
Received: November 6, 2001

Dear Mr. Zott:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

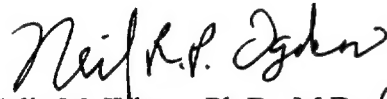
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.


Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4659. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



Celia M. Witten, Ph.D., M.D. 
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

V. INDICATIONS FOR USE

510(k) Number (if known): K013671

Device Name: L-25 Absorbable Surgical Suture (Trade name to be determined)

Indications For Use:

L-25 synthetic absorbable sutures are indicated for use in general soft tissue approximation and/or ligation, but not for use in cardiovascular or neurological surgery, microsurgery, or ophthalmic surgery.

(Please do not write below this line - continue on another page if needed)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use: X OR Over-The-Counter Use: _____
(Per 21 CFR 801.109)

NLD for cmw
(Division Sign-Off)
Division of General, Restorative
and Neurological Devices

510(k) Number K013671